

(19) JAPANESE PATENT OFFICE (JP)

(12) Official Gazette for Unexamined Patent Applications (A)

(11) Japanese Unexamined Patent Application
(Kokai) No. 2001-183502

(P2001-183502A)

(43) Disclosure Date: 6 July 2001 (2001.6.7)

(51)	Int.Cl. ⁷	Ident. Symbols	FI	Topic Code (Reference)
	G02B	1/04	G02B	1/04
	C08F	290/06	C08F	290/06
		299/08		299/08
	G02C	13/00	G02C	13/00
Request for Examination: Not yet requested				Number of Claims: 3 OL (Total of 13 pages)
(21) Application No.:	Hei 11-367752			
(22) Application Date:	24 December 1999 (1999.12.24)			
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(continued on last page)				

(54) [Title of the Invention] Soft Contact Lens Materials Resistance to Soiling Attachment

(57) Abstract

[Problem] To provide a water-containing soft contact lens material to which little soiling is attached and which has a superior oxygen permeation coefficient (DK value), optical and mechanical properties.

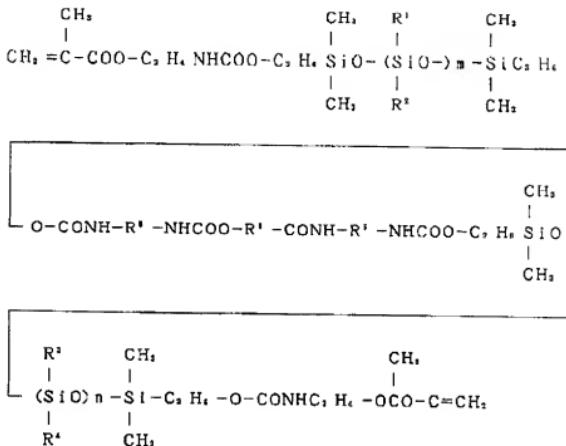
[Means of Solution] A contact lens material that is comprised of a polymer that is obtained by polymerizing one or two or more bifunctional organic siloxane macromers that are represented by a specified general formula with said bifunctional organic siloxane macromers only or together with one or two or more monomers that can be copolymerized.

[Claims]

[Claim 1] A contact lens material that is comprised of a polymer that is obtained by polymerizing one or two or more bifunctional organic siloxane macromers that are represented by chemical formula 1 below with said bifunctional organic

siloxane macromers only or together with one or two or more monomers that can be copolymerized.

[Chemical Formula 1]

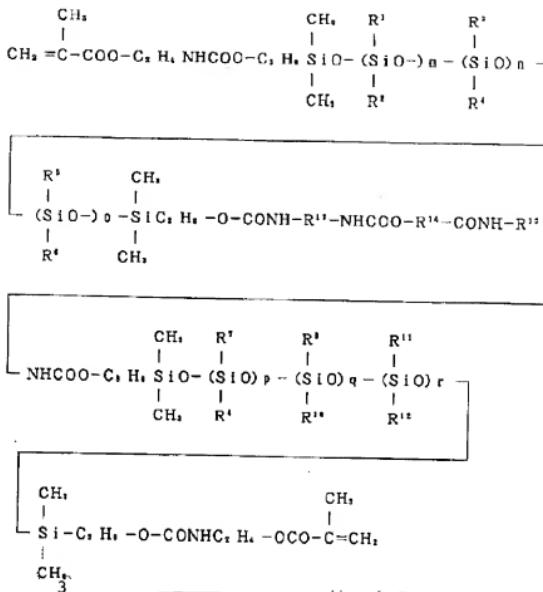


[Wherein, R¹, R², R³ and R⁴ are groups selected from -CH₃ and -CH₂CH₂CF₃ and may be the same or different, m and n are integers in the range of 10 to 100, R⁵ is a group selected from hexamethylene groups, tetramethylene groups, dicyclohexylmethane groups, hydrogenated tolylene groups, hydrogenated xylylene groups and isophorone groups, R⁶ is a group selected from -(C₂H₄O)₁₋₁₀, -(CH₂CHO)₁₋₁₀ and -(C₄H₈O)₁₋₁₀]

and 1 is an integer in the range of 1 to 20.

[Claim 2] A contact lens material that is comprised of a polymer that is obtained by polymerizing one or two or more bifunctional organic siloxane macromers that are represented by chemical formula 2 below with said

bifunctional organic siloxane macromers only or together with one or two or more monomers that can be copolymerized.



[Wherein, R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R¹¹ and R¹² are groups selected from -CH₃ and -CH₂CH₂CF₃ and may be the same or different. m, n, o, p and r are integers in the range of 5 to 30. R³, R⁴, R⁵ and R¹⁰ are selected from -CH₃,

-(CH₂CH₂O)sCH₂ and may be the same or different. n and q are integers in the range of 5 to 30. s is an integer in the range of 1 to 20. R¹¹ is a group selected from -(C₂H₄O)_l, -(CH₂CHO)_l -

range of 1 to 20. R¹³ is a group selected from hexamethylene groups, tetramethylene groups, cyclohexylmethane groups, hydrogenated tolylene groups, hydrogenated xylylene groups and isophorone groups.

and -(C₄H₈O)_l- and l is an integer in the range of 1 to 20.

[Claim 3] A contact lens material as described in Claim 1 or 2 in which the copolymerizable monomer is a methacrylic acid siloxanyl alkyl ester, an acrylic acid fluoroalkyl ester, a methacrylic acid fluoroalkyl ester, an acrylic acid alkyl ester and a methacrylic acid alkyl ester, an acrylic acid alkyl ester and a methacrylic acid alkyl ester of an alkyl group having an hydroxyl group, an N,N'-dimethylacrylamide, N-vinyl pyrrolidone, ethylene glycol dimethacrylate and tetraethylene dimethacrylate.

[Detailed Description of the Invention]

[0001]

[Technological field of the invention] This invention relates to a novel contact lens material. Speaking in greater detail, this invention relates to a soft contact lens material which exhibits little attachment of soiling such as the proteins in tears and which exhibits a superior balance of physical properties such as oxygen permeability.

[0002]

[Prior art] Soft contact lenses can be divided into water-containing soft contact lenses and water non-containing soft contact lenses. Water-containing soft contact lenses are comprised of copolymers that are obtained by polymerizing

hydrophilic monomers such as hydroxyethyl methacrylate and N-vinyl pyrrolidone as the principal constituents. The lenses are made by the cutting or casting methods and the lenses, which have been treated with physiological saline solution have a water content ratio of approximately 40 to 70%.

[0003] Known non-water containing soft contact lenses include, for example, silicone rubber lenses that are obtained by a method in which a platinum catalyst is added to a mixture of polydimethylene siloxane and methyl hydrodiene polysiloxane of which both terminals of the molecular chain are blocked by vinyl dimethyl silyl groups and the mixture is heated and hardened by the molding method and flexible lenses of which the principal constituent is polyperfluoroether to the two terminals to which have been added polymerizable groups such as methacryloxy groups and which have a modulus of elasticity intermediate between those of soft contact lenses and hard contact lenses (Japanese Patent Application Early Disclosure No. Sho 54-81363 [1979] and Japanese Patent Application Early Disclosure No. Sho 58-127914 [1983]).

[0004] In addition, non-water-containing soft contact lenses of excellent wearing feel can also be manufactured by cutting from a hard substrate in which (meth) acrylic acid and an acrylic acid ester have been copolymerized to make a lens, which lens is then subjected to esterification or ester interchange treatment (Japanese Patent Application Early Disclosure No. Sho 48-75047 [1973]). Contact lens materials of high oxygen permeability of which polyfunctional polysiloxanes are the principal constituents have been disclosed in Japanese Patent Application Early Disclosure No. Sho 54-24047 [1979], Japanese Patent Application Early Disclosure No. Sho 56-51715 [1981], Japanese Patent Application Early Disclosure No. Sho 59-229524 [1984] and Japanese Patent Application Early Disclosure No. Hei 2-188717 [1990]. Water-containing soft contact lenses have the advantages that they are soft and of excellent wearing feel and that they have high oxygen permeability when they are of a high water content type. However, they are easily damaged, are of poor durability, are easily contaminated by components of tears and there is the danger of growth of bacteria, for which reason they must be periodically boiled and disinfected and there are many inconveniences in handling. The following problems are seen with non-water-containing contact lenses.

[0005] First, silicone rubber contact lenses have the drawback that they adhere to the cornea because the layer that had been subjected to hydrophilic treatment, which is performed on initial development to improve the hydrophobicity of the lens surface, is peeled off and its elasticity becomes excessively great. They have not yet reached the level of widespread practical use. Although most recently there has been some improvement in the above

described drawbacks as the result of advances in surface hydrophilic treatment, changes in lens design and improvements in mechanical properties, they are still not sufficiently satisfactory and have not reached the point of widespread practical use.

[0006] Next, because non-water-containing lenses of which polyperfluoroether is the principal constituent are closer to the semi-hard type than soft contact lenses, a wearing feel like that of soft contact lenses is not obtained and movement of the lens on the cornea is not satisfactory. In addition, soft contact lenses of which (meth) acrylic acid esters are the principal constituents have poor oxygen permeability and are not suited to wearing for long periods. Materials of which polyfunctional polysiloxanes are the principal constituents are materials that are close to silicone rubber lenses, not taking into consideration contamination by components of tears.

[0007]

[Problems the invention is intended to solve] This invention has the objective of providing, on the basis of these facts, a water-containing soft contact lens with which there is little soiling attachment and that has a superior oxygen transmission coefficient (DK value) and optical and mechanical properties.

[0008]

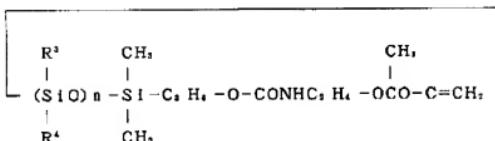
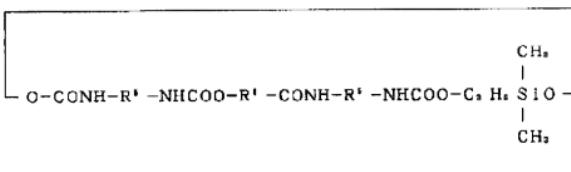
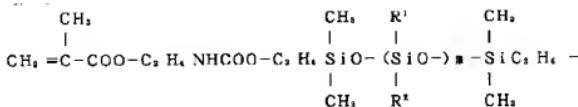
[Means for solving the problems] The inventors conducted repeated and intensive research for the purpose of developing a soft contact lens material with which the above-described problems could be solved. As the result, they discovered that this objective could be achieved by means of a material comprised of copolymers obtained by polymerizing bifunctional organic siloxane macromers having a specified structure and molecular weight as the principal constituents and perfected this invention on the basis of this finding.

[0009] Specifically, this invention is as described below.

1. A contact lens material that is comprised of a polymer that is obtained by polymerizing one or two or more bifunctional organic siloxane macromers that are represented by chemical formula 1 below with said bifunctional organic siloxane macromers only or together with one or two or more monomer that can be copolymerized.

[0010]

[Chemical Formula 1]



[Wherein, R^1 , R^2 , R^3 and R^4 are groups selected from $-\text{CH}_3$ and $-\text{CH}_2\text{CH}_2\text{CF}_3$ and may be the same or different, m and n are integers in the range of 10 to 100. R^5 is a group selected from hexamethylene groups, tetramethylene

groups, dicyclohexylmethane groups, hydrogenated tolylene groups, hydrogenated xylylene groups and isophorone groups.

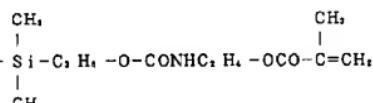
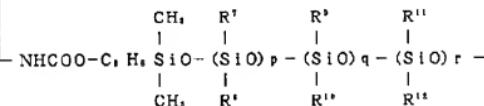
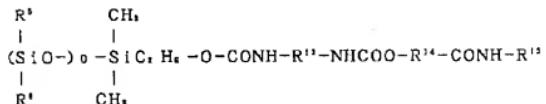
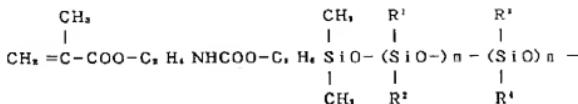
R^6 is a group selected from $-(\text{C}_2\text{H}_4\text{O})_l$, $-(\text{CH}_2\text{CHO})_l$ and $-(\text{C}_4\text{H}_8\text{O})_l$



and l is an integer in the range of 1 to 20.

2. A contact lens material that is comprised of a polymer that is obtained by polymerizing one or two or more bifunctional organic siloxane macromers that are represented by chemical formula 2 below with said bifunctional organic siloxane macromers only or together with one or two or more monomers that can be copolymerized.

[Chemical Formula 2]



- 3 -

[Wherein, R^1 , R^2 , R^5 , R^6 , R^7 , R^8 , R^{11} and R^{12} are groups selected from $-\text{CH}_3$ and $-\text{CH}_2\text{CH}_2\text{CF}_3$ and may be the same or different. m , o , p and r are integers in the range of 5 to 30. R^3 , R^4 , R^9 and R^{10} are selected from $-\text{CH}_3$,

$-(\text{CH}_2\text{CH}_2\text{O})_s\text{CH}_2$ and may be the same or different, n and q are integers in the range of 5 to 30, s is an integer in the range of 1 to 20, R^{13} is a group selected from hexamethylene groups, tetramethylene groups, cyclohexylmethane groups, hydrogenated tolylene groups, hydrogenated xylylene groups and isophorone groups, R^{14} is a group selected from $-(\text{C}_2\text{H}_4\text{O})_l$, $-(\text{CH}_2\text{CHO})_l$ and

$-(\text{C}_2\text{H}_5\text{O})_1 - \text{CH}_3$, and l is an integer in the range of 1 to 20.]

[0012] The R^5 group in Chemical Formula 1 and the R^{13} group in Chemical Formula 2 are residues of bifunctional isocyanates and may include, for example, residues of hexamethylene diisocyanate, tetramethylene diisocyanate, 2,4,4-trimethyl-1,6-hexamethylene diisocyanate, 2,6-diisocyanatomethyl caproate, 3-isocyanatomethyl-3,5,5-trimethyl cyclohexyl isocyanate, dicyclohexylmethane-4,4'-diisocyanate, hydrogenated xylylene diisocyanate, hydrogenated tolylene diisocyanate, and isophorone

diisocyanate, of which residues of hexamethylene diisocyanate and isophorone diisocyanate are desirable in terms of the balance of the mechanical properties of the lens. The R^6 group and the R^{14} group are residues of polyoxyalylene glycols, and include, for example, residues of polyethylene glycol, polypropylene glycol and polytetraethylene glycol. Substances of a degree of polymerization in the range of 10 to 50 can be used. Of these, polyethylene glycol is preferable because of its superior aqueous wetting properties and properties of preventing soiling attachment.

[0013] The bifunctional organic siloxane macromers represented by Chemical Formula 1 and Chemical Formula 2 are obtained, for example, by reacting a modified terminal isocyanate of polyethylene glycol obtained by reacting twice the volume of a diol having a polysiloxane structure, for example, by reacting hexamethylene isocyanate with twice its volume of polyethylene glycol, which diol is obtained by a ring-opening insertion reaction of 1,3-bis-(hydroxypropyl) tetramethyl disiloxane with octamethylcyclotetrasiloxane and 1,3,5-trifluoropropyl trimethyl cyclotrisiloxane. The structures of the bifunctional organic siloxane macromers that have been

obtained in this way were analyzed from their NMR (nuclear magnetic resonance spectra) and IR absorption spectra. The structural ratios of the dimethyl siloxane structure and the methyl trifluoropropyl siloxane structure of the polysiloxane structure were found by determining the integration value of proton absorption relevant to the methyl groups and of proton absorption relevant to the trifluoropropyl groups that appeared on the NMR spectra and by determining the integration values of proton absorption relevant to the methylene groups contained in the same molecules.

[0014] The contact lenses of this invention are comprised of polymers obtained by copolymerizing one or two or more of the bifunctional organic siloxane macromers represented by Chemical Formula 1 and Chemical Formula 2 only with said bifunctional organic siloxane macromers or with one or two or more copolymerizable monomers. The type and binding number of the structural units that constitute the R⁶ group and the R¹⁴ group and the residues that constitute the R⁵ group and the R¹³ group in chemical formula 1 and chemical formula 2 depend on the target properties of contact lens, for example, the required oxygen permeability, softness, difficulty of soiling attachment and strength.

[0015] We shall now explain the copolymerizable monomers. The methacrylic acid siloxanyl alkyl esters that are used as the copolymerizable monomers in this invention are the above described siloxane macromers and substances that improve the oxygen permeability of the lens and have the effects of aiding the compatibility of the monomers and macromers described below and of increasing the light ray transmission ratio of the lens. For example, one or two or more can be selected from tris(trimethylsiloxy) silyl propyl methacrylate, bis(trimethylsiloxy)methyl silyl propyl methacrylate, pentamethyl disiloxane propyl methacrylate, tris(trimethylsiloxy) silyl propyloxyethyl methacrylate, and tris(polydimethylsiloxy) silyl propyl methacrylate.

[0016] The acrylic acid fluoroalkyl esters and methacrylic fluoroalkyl esters that are used as the copolymerizable monomers in this invention have the properties of water repellency and oil repellency due to decrease in critical surface tension attributable to their fluorine atoms. These properties have the effect of inhibiting contamination of the surface of the contact lens by such constituents of tears as proteins and lipids. In terms of physical properties, it is further desirable that the glass transition point of the monomer constituent be below room temperature so that the softness and flexibility required of soft contact lenses when it is copolymerized will be manifested.

[0017] Specific examples of these acrylic acid fluoroalkyl esters and methacrylic fluoroalkyl esters can include trifluoroethyl acrylate, tetrafluoroethyl acrylate, tetrafluoropropyl acrylate, pentafluoropropyl acrylate, hexafluorobutyl acrylate, hexafluoroisopropyl acrylate, heptafluorobutyl acrylate, octafluoropentyl acrylate, nonafluoropentyl acrylate, dodecafluoropentyl acrylate, dodecafluorobutyl acrylate, dodecafluoroctyl

acrylatetradecalfluoroheptyl acrylate and methacrylate corresponding to these acrylates, and, preferably, with trifluoroethyl acrylate, hexafluoroisopropyl acrylate, octafluoropentylacrylate and dodecafluoroctyl acrylate being used. One of these monomers may be used and combinations of two or more may also be used.

[0018] In addition, the acrylic acid alkyl ester monomers and methacrylic acid alkyl ester monomers that are used as copolymerizable monomers in this invention have the effect of improving the compatibility of the bifunctional organic siloxane macromers and the (meth) acrylic acid fluoroalkyl esters and play a role in greatly broadening the range of use. Preferably, it is desirable that they be substances that lower the glass transition point of the copolymer to room temperature or to 0°C or lower. Thereby, they have the effect of conferring elasticity and softness on the copolymer.

[0019] Specific examples of these acrylic acid alkyl esters and methacrylic acid alkyl esters can include methyl acrylate, ethyl acrylate, n-propyl acrylate, n-butyl acrylate, tert-butyl acrylate, isobutyl acrylate, n-hexyl acrylate, n-octyl acrylate, n-heptyl acrylate, n-nonyl acrylate, n-decyl acrylate, isodecyl acrylate, n-lauryl acrylate, tridecyl acrylate, n-dodecyl acrylate, cyclopentyl acrylate, cyclohexyl acrylate, n-acetyl acrylate and methacrylates corresponding to these acrylates. Preferably, n-butyl acrylate, n-octyl acrylate, n-lauryl methacrylate and n-stearyl methacrylate are used. One of these monomers may be used or two or more may be used in combination. As desired, the monomers indicated below can be copolymerized for the purpose of improving mechanical properties, surface wettability and the dimensional stability of the lens.

[0020] Monomers for improving mechanical properties include, for example, aromatic vinyl compounds such as styrene, tert-butyl styrene and α -methyl styrene. Monomers for improving surface wettability include, for example, methacrylic acid, acrylic acid, itaconic acid, 2-hydroxyethyl methacrylate, 2-hydroxyethyl acrylate, 2-hydroxypropyl methacrylate, 2-hydroxypropyl acrylate, glycerol methacrylate, polyethylene glycol methacrylate, N,N'-dimethyl acrylamide, N-methyl acrylamide, dimethyl aminoethyl methacrylate, methylene bisacrylamide, diacetone acrylamide and N-vinyl pyrrolidone.

[0021] Monomers for improving the dimensional stability of the lens can include, for example, ethylene glycol dimethacrylate, diethylene glycol dimethacrylate, triethylene glycol dimethacrylate, polyethylene glycol dimethacrylate, trimethylol propane trimethacrylate, pentaerythritol trimethacrylate, acryl methacrylate and acrylates corresponding to these methacrylates. One of these monomers may be used or two or more of them may be used in combination. Mixed monomers are employed in which copolymerizable monomers are combined for the purpose of improving the optical properties, oxygen permeability, mechanical strength, deformation recovery capacity, soiling attachment when worn in the eyes,

dimensional stability in tears and the balance of these properties of the contact lens in changes over time.

[0022] Examples of mixed monomers that are copolymerizable with the bifunctional organic siloxane macromers of this invention include combinations of tris (trimethylsiloxy) silyl propyl methacrylate, trifluoroethyl methacrylate, N-vinyl pyrrolidone, N,N'-dimethyl acrylamide and ethylene glycol dimethacrylate, and combinations of N-vinyl pyrrolidone, N,N'-dimethyl acrylamide and ethylene glycol dimethacrylate.

[0023] The copolymers that are used as the contact lens materials of this invention can be molded into contact lenses by filling a mold with the monomer mixture and by known methods, including the cast polymerization method in which it is subjected to radical polymerization, a method in which the monomer mixture is charged into a rotating hemispherical mold and is polymerized, or a method in which the copolymer is frozen and cut at a low temperature. The copolymerization methods that can be used include a method in which a photopolymerization initiator such as benzoin, benzophenol or benzyl dimethyl ketanol is present in the monomer mixture and polymerization is effected by irradiation with ultraviolet rays or a method in which thermal polymerization is performed using azo compounds and organic such as azobisisobutyronitrile, benzoyl peroxide and lauroyl peroxide.

[0024]

[Working Examples] Next, we shall present a more detailed explanation of this invention by means of working examples. However, this invention is not limited in any way by these examples. The various physical properties were found as follows.

(1) Oxygen permeability coefficient

A K-316 · IPI gas permeability determination device manufactured by the Rika Seiki Industrial Company (Ltd.) was used. A test material strip in the shape of a lens of a diameter of 13.5 mm and a thickness of 0.09 mm was used for the determination and the oxygen permeability coefficient was calculated from the slope of the straight line on the chart. Determinations were performed at 35°C. The numerical value was indicated by $\times 10^{-11}$ (ml · cm/cm² · sec · mmHg).

[0025] (2) Contact angle

A "Contact Angle Meter Model CA-DT" manufactured by the Kyowa Surface Science Company (Ltd.) was used. The test material strips were used in the determinations after having been immersed for 24 hours in purified water before the determinations. In the determination, the test material strip was allowed to stand in purified water and air bubbles of 1.5 to 2.0 mm in diameter were brought into contact with the bottom face of the test material strip by means of a syringe. The angle of contact that the air bubbles and the test material strip formed was read and the angle of contact

was found by calculation. Determinations were performed at 24°C.

(3) Tensile strength

An SV-50 universal tensile compression testing machine manufactured by Shimazu Seisakujo was used. The test material strips were supplied for the determinations as samples cut to a width of 5.0 mm. The drawing speed was 10 mm/minute. Three determinations were performed for the same test material strip and tensile strength was found from the values that were obtained by the calculation method described in JIS-K-6301.

[0026] (4) Soiling attachment capacity

Attachment capacity of proteins

A model soiling aqueous solution having the composition described below was mixed.

NaCl	0.9 wt%
NaH ₂ PO ₄	0.0184 wt%
Na ₂ HPO ₄	0.076 wt%
Egg white lysozyme	0.12 wt%
Bovine serum albumin	0.388 wt%
CaCl ₂ (H ₂ O) ₂	0.0407 wt%

One lens was introduced into 2 ml of this solution and was soaked in it for 24 hours at a temperature of 37°C. The lens was then introduced into purified water and was shaken and washed for 30 minutes. The lens was removed and the surface water was gently wiped off. The lens was immersed in a protein analysis reagent (BCA solution) solution (1 lens/2 ml) and a reaction was carried out for 5 hours at 40°C, after which determinations were made at UV 562 nm with a spectrophotometer (Model V-550, manufacture by Nippon Bunko Co. (Ltd.)). The mass of attached protein per lens was found from a calibration curve that had been found separately.

[0027] (5) Attachment capacity of lipids

A 1% phosphate buffer solution of olive oil was stirred and mixed, the lens was introduced and was immersed in it for 23 hours in a constant temperature tank at 40°C. The lens was then introduced into a vessel into which 5 ml of purified water had been introduced and shaking and washing were carried out for 30 seconds. This washing was repeated 5 times. The lens was vacuum dried, after which extraction was performed with a mixed solvent of chloroform/methanol : 2/1, coloration was effected with triglyceride G test solution and absorbance was determined with a spectrophotometer (Model V-550, manufacture by Nippon Bunko Co. (Ltd.)). The quantity of attached lipid per lens was found from a calibration curve that has been found separately.

[0028] (6) Water content ratio

The test material was immersed for 72 hours in purified water at 37°C, the water on the surface was gently wiped

off and weight was precisely measured, after which it was dried in a vacuum dryer until a constant weight was reached.

[0029]

[Working Example 1] 100 g (0.5 mol) of polyethylene glycol of molecular weight 200 was charged into a 500 ml brown, round-bottom flask equipped with a condenser, a thermometer and a stirrer, 201.6 g (1.2 mol) of hexamethylene diisocyanate was added, 5 drops of dibutyltin dilaurate and tetrahydrofuran were added, and, as the mixture was being stirred in a nitrogen atmosphere, a reaction was carried out for 5 hours at 70°C. During the reaction, changes in absorption were followed by means of the infrared absorption spectrum on the basis of the isocyanate groups and elimination in changes in absorption was confirmed.

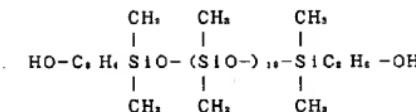
[0030] 313.3 g (0.2 mol) of organic siloxane diol as represented by Chemical Formula 3 below and having hydroxyl groups in both terminals was charged into a 500 ml brown, round-bottom flask equipped with a condenser, a

thermometer and a stirrer, 53.6 g (0.01 mol) of the above described diisocyanate compound was added, 5 drops of dibutyltin dilaurate and tetrahydrofuran were added, and as the mixture was being stirred in a nitrogen atmosphere, a reaction was carried out for 5 hours at 70°C. During the reaction, changes in absorption were followed by means of the infrared absorption spectrum on the basis of the isocyanate groups and elimination in changes in absorption was confirmed. In addition, 37.2 g (0.24 mol) of 2-isocyanatoethyl methacrylate was added dropwise, the reaction was continued and changes in absorption were followed by means of the infrared absorption spectrum on the basis of the isocyanate groups and elimination in changes in absorption was confirmed.

[0031]

[Chemical Formula 5]

[Chemical Formula 3]



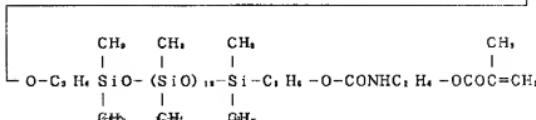
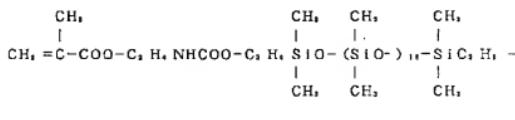
After cooling, approximately 300 ml of tetrahydrofuran was added, activated charcoal for decolorization and magnesium sulfate for dehydration were added and the mixture was allowed to stand overnight. Following that, the mixture was filtered with an 0.2 μ Teflon filter, after which the low boiling point substances such as tetrahydrofuran were removed using an evaporator. When this was done, 21.6 g of a transparent, slightly viscous

liquid was obtained. The structure of this liquid was analyzed by NMR (proton nuclear magnetic resonance spectrum). As the result, it was confirmed that it was an organic siloxane macromer as represented by Chemical formula 4 below.

[0032]

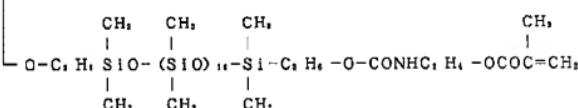
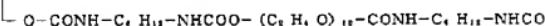
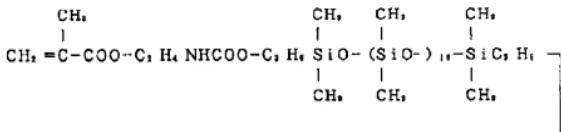
[Chemical Formula 6]

[Chemical Formula 4]



55 parts by weight of the organic siloxane macromer containing urethane bonds indicated by Chemical Formula 4, 15 parts by weight of tris(trimethylsiloxy) silyl propyl methacrylate, 15 parts by weight N,N-dimethyl acrylamide, 5 parts by weight of trifluoroethyl methacrylate, 1 part by weight of ethylene glycol dimethacrylate and 0.5 parts by weight of benzyl dimethyl ketanol (brand name: I-651), manufactured by the Ciba-Geigy Company) were added, and the mixture was dissolved and mixed for approximately 1 hour in a nitrogen atmosphere with a magnetic stirrer. Following that, the above-described reaction solution was poured into a polypropylene mold cell for the lens of a base cover of 9.0 mm, a size of 13.5 mm, a power of -3.00 diopters and a central thickness of 0.09 mm, said cell was irradiated for approximately 15 minutes with ultraviolet rays at a temperature of 40 to 50°C and a transparent copolymer was obtained.

[Chemical Formula 5]



Organic siloxane diol was reacted with 2 times its volume of 2-isocyanatoethyl methacrylate in the same way as in Working Example 1 and the mixture was purified, with the organic siloxane containing urethane bonds as indicated by Chemical Formula 5 being obtained. 55 parts by weight of the organic siloxane macromer containing urethane bonds as indicated by the above-described Chemical Formula 5, 15 parts by weight of tris(trimethylsiloxy) silyl propyl methacrylate, 15 parts by weight of N,N-dimethyl acrylamide, 5 parts by weight of trifluoroethyl methacrylate, 1 part by weight of ethylene glycol dimethacrylate and 0.5 parts by weight of benzyl dimethyl ketanol (brand name: I-651, manufactured by the Ciba-Geigy Company) were added, and the mixture was dissolved and mixed for approximately 1 hour in a nitrogen atmosphere with a magnetic stirrer. Following that, the above-described reaction solution was poured into a polypropylene mold cell for the lens of a base cover of 9.0 mm, a size of 13.5 mm, a power of -3.00 diopters and a central thickness of 0.09 mm, said cell was irradiated for

[0033] The copolymer that was obtained in this way was immersed and swollen in purified water at 90°C and was then further swollen in purified water at 70°C, with a water containing soft lens being obtained. In addition, a lens-shaped water-containing substance of a power of 0.0 diopters was obtained in the same way. The water-containing ratio was 28%, the oxygen permeability coefficient was 156, the contact angle was 26°, tensile strength was 216 g/mm², and protein and lipid soiling attachment were 12 µg/lens and 40 µg/lens.

[0034]

[Working Example 2]

[0035]

[Chemical Formula 7]

approximately 15 minutes with ultraviolet rays at a temperature of 40 to 50°C and a transparent copolymer was obtained.

[0036] The copolymer that was obtained in this way was immersed and swollen in purified water at 90°C and was then further swollen in purified water at 70°C, with a water containing soft lens being obtained. In addition, a lens-shaped water-containing substance of a power of 0.0 diopters was obtained in the same way. The water-containing ratio was 30%, the oxygen permeability coefficient was 150, the contact angle was 22°, tensile strength was 216 g/mm², and protein and lipid soiling attachment were 10 µg/lens and 40 µg/lens.

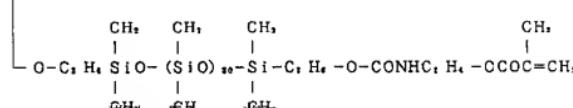
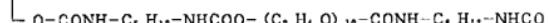
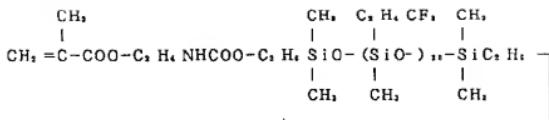
[0037]

[Working Example 3]

[0038]

[Chemical Formula 8]

[Chemical Formula 6]



Organic siloxane diol was reacted with 2 times its volume of 2-isocyanatoethyl methacrylate in the same way as in Working Example 1 and the mixture was purified, with the organic siloxane containing urethane bonds as indicated by Chemical Formula 6 being obtained. 55 parts by weight of the organic siloxane macromer containing urethane bonds as indicated by the above-described Chemical Formula 6, 15 parts by weight of tris(trimethylsiloxy) silyl propyl methacrylate, 15 parts by weight of N-vinyl pyrrolidone, 15 parts by weight of N,N'-dimethyl acrylamide, 5 parts by weight of trifluoroethyl methacrylate, 1 part by weight of ethylene glycol dimethacrylate and 0.5 parts by weight of benzyl dimethyl ketanol (brand name: I-651, manufactured by the Ciba-Geigy Company) were added, and the mixture was dissolved and mixed for approximately 1 hour in a nitrogen atmosphere with a magnetic stirrer. Following that, the above-described reaction solution was poured into a polypropylene mold cell for the lens of a base cover of 9.0 mm, a size of 13.5 mm, a power of -3.00 diopters and a central thickness of 0.09 mm, said cell was irradiated for

approximately 15 minutes with ultraviolet rays at a temperature of 40 to 50°C and a transparent copolymer was obtained.

[0039] The copolymer that was obtained in this way was immersed and swollen in purified water at 90°C and was then further swollen in purified water at 70°C, with a water containing soft lens being obtained. In addition, a lens-shaped water-containing substance of a power of 0.0 diopters was obtained in the same way. The water-containing ratio was 30%, the oxygen permeability coefficient was 160, the contact angle was 22°, tensile strength was 216 g/mm², and protein and lipid soiling attachment were 10 µg/lens and 40 µg/lens.

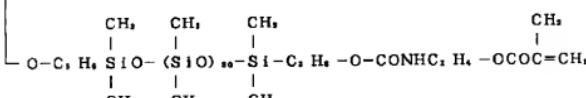
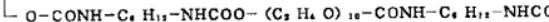
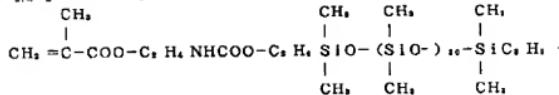
[0040]

[Working Example 4]

[0041]

[Chemical Formula 9]

[Chemical Formula 7]



Organic siloxane diol was reacted with 2 times its volume of 2-isocyanatoethyl methacrylate in the same way as in Working Example 1 and the mixture was purified, with the organic siloxane containing urethane bonds as indicated by Chemical Formula 7 being obtained. 55 parts by weight of the organic siloxane macromer containing urethane bonds as indicated by the above-described Chemical Formula 7, 15 parts by weight of tris(trimethylsiloxy) silyl propyl methacrylate, 15 parts by weight of N-vinyl pyrrolidone, 15 parts by weight of N,N'-dimethyl acrylamide, 5 parts by weight of trifluoroethyl methacrylate, 1 part by weight of ethylene glycol dimethacrylate and 0.5 parts by weight of benzyl dimethyl ketanol (brand name: I-651, manufactured by the Ciba-Geigy Company) were added, and the mixture was dissolved and mixed for approximately 1 hour in a nitrogen atmosphere with a magnetic stirrer. Following that, the above-described reaction solution was poured into a polypropylene mold cell for the lens of a base cover of 9.0 mm, a size of 13.5 mm, a power of -3.00 diopters and a central thickness of 0.09 mm, said cell was irradiated for

approximately 15 minutes with ultraviolet rays at a temperature of 40 to 50°C and a transparent copolymer was obtained.

[0042] The copolymer that was obtained in this way was immersed and swollen in purified water at 90°C and was then further swollen in purified water at 70°C, with a water containing soft lens being obtained. In addition, a lens-shaped water-containing substance of a power of 0.0 diopters was obtained in the same way. The water-containing ratio was 30%, the oxygen permeability coefficient was 210, the contact angle was 23°, tensile strength was 156 g/mm², and protein and lipid soiling attachment were 15 µg/lens and 44 µg/lens.

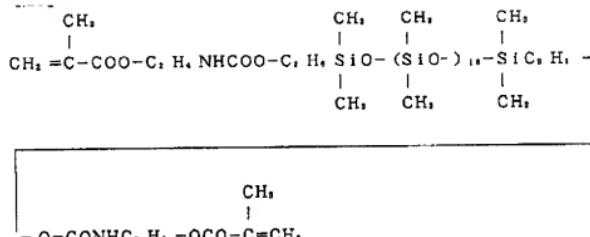
[0043]

[Comparative Example 1]

[0044]

[Chemical Formula 10]

[Chemical Formula 8]



Organic siloxane diol was reacted with 2 times its volume of 2-isocyanatoethyl methacrylate in the same way as in Working Example 1 and the mixture was purified, with the organic siloxane containing urethane bonds as indicated by Chemical Formula 8 being obtained. 55 parts by weight of the organic siloxane macromer containing urethane bonds as indicated by the above-described Chemical Formula 8, 15 parts by weight of tris(trimethylsiloxy) silyl propyl methacrylate, 15 parts by weight of N-vinyl pyrrolidone, 15 parts by weight of N,N'-dimethyl acrylamide, 5 parts by weight of trifluoroethyl methacrylate, 1 part by weight of ethylene glycol dimethacrylate and 0.5 parts by weight of benzyl dimethyl ketanol (brand name: I-651, manufactured by the Ciba-Geigy Company) were added, and the mixture was dissolved and mixed for approximately 1 hour in a nitrogen atmosphere with a magnetic stirrer. Following that, the above-described reaction solution was poured into a polypropylene mold cell for the lens of a base cover of 9.0 mm, a size of 13.5 mm, a power of -3.00 diopters and a central thickness of 0.09 mm, said cell was irradiated for approximately 15 minutes with ultraviolet rays at a temperature of 40 to 50°C and a transparent copolymer was obtained.

[0045] The copolymer that was obtained in this way was immersed and swollen in purified water at 90°C and was then further swollen in purified water at 70°C, with a water containing soft lens being obtained. In addition, a lens-shaped water-containing substance of a power of 0.0 diopters was obtained in the same way. The water-containing ratio was 25%, the oxygen permeability coefficient was 250, the contact angle was 27°, tensile strength was 156 g/mm², and protein and lipid soiling attachment were 30 µg/lens and 35 µg/lens. This lens was installed for 21 days in the eyes of a domestic rabbit. There was a large amount of lipid soiling, the lens exhibited white turbidity and the wettability of the recovered lens was poor. When the lens was removed from the storage solution, there was no aqueous phase on the lens surface. In addition, the lens was greatly swollen and absorption of lipids was found.

[0046]

[Effect of the invention] The contact lens material of this invention is a material of superior soiling attachment properties and of a superior balance of physical properties such as oxygen permeability and mechanical strength.

Continued from front page

F Terms (Reference)	2H006	BB01	BB05	BB07		
	4J027	AC03	AC06	AG04	AG06	AG09
		AG24	AG27	BA05	BA06	BA07
		BA08	BA14	BA15	BA17	BA19
		BA20	BA26	BA27	CB04	CB10
		CC02	CC05	CD04		